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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (currently amended) A recombinant virus <u>baculovirus</u> eapable of infecting a nonpermissive cell, comprising:

a first nucleic acid sequence encoding a detectable marker operably linked to a first promoter, wherein the first promoter is active in a <u>permissive</u> host cell and inactive in a non-permissive cell; and

a second nucleic acid sequence which includes an exogenous nucleic acid sequence operably linked to a second promoter, wherein the second promoter is active in the non-permissive cell and inactive in the permissive cell.

2. (canceled)

- 3. (currently amended) The recombinant $\frac{1}{2}$, wherein the first promoter is inactive and the second promoter is active in a mammalian cell.
- 4. (currently amended) The recombinant $\frac{1}{2}$, wherein the first promoter is inactive and the second promoter is active in a human cell.
- 5. (currently amended) The recombinant $\frac{1}{2}$, wherein the first promoter is inactive and the second promoter is active in a primary human cell.
- 6. (currently amended) The recombinant virus baculovirus of claim 1 2, wherein the first promoter is inactive and the second promoter is active in a non-permissive insect cell.

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7. (currently amended) The recombinant <u>virus</u> <u>baculovirus</u> of claim 6, wherein the first promoter is inactive and the second promoter is active in a non-permissive *Drosophila* cell.

- 8. (currently amended) The <u>virus</u> <u>baculovirus</u> of claim 1, wherein the first promoter is a viral polyhedrin promoter.
- 9. (currently amended) The recombinant virus <u>baculovirus</u> of claim 1, wherein the first promoter is a P10 promoter.
- 10. (currently amended) The recombinant virus <u>baculovirus</u> of claim 3, wherein the second promoter is a CMV promoter, an RSV promoter, or an SV40 promoter.
- 11. (currently amended) The recombinant virus <u>baculovirus</u> of claim 6, wherein the second promoter is a heat shock protein promoter, an Orgyia pseudotsugata immediate-early promoter, a metallothionein (MT) promoter, or an actin 5C promoter.
- 12. (currently amended) The recombinant virus <u>baculovirus</u> of claim 1, wherein the detectable marker is a fluorescent protein.
- 13. (currently amended) The recombinant virus baculovirus of claim 12, wherein the fluorescent protein is green fluorescent protein (GFP), enhanced GFP (EGFP), enhanced yellow fluorescent protein (EYFP), enhanced cyan fluorescent protein (ECFP), enhanced blue fluorescent protein (EBFP), or <u>Discosoma</u> red fluorescent protein (DsRed).
- 14. (currently amended) A method for selecting a viral plaque for infection of non-permissive cells, comprising:

providing a recombinant virus capable of infecting a non-permissive cell, which virus includes baculovirus that comprises a first nucleic acid sequence encoding a detectable marker operably linked to a first promoter, wherein the first promoter is active in a permissive host cell

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culture and is inactive in the <u>a</u> non-permissive cell; and a second nucleic acid sequence comprising an exogenous nucleic acid sequence operably linked to a second promoter, wherein the second promoter is active in the non-permissive cell <u>and inactive</u> in the permissive cell;

infecting a the permissive host cell culture with the recombinant baculovirus; and identifying a viral plaque by detecting expression of the detectable marker, thereby selecting a viral plaque for infection of non-permissive cells.

15. (canceled)

- 16. (currently amended) The method of claim 14 15, wherein a recombinant virus baculovirus is provided in which the first promoter is inactive and the second promoter is active in a mammalian cell.
- 17. (currently amended) The method of claim 16, wherein a recombinant virus <u>baculovirus</u> is provided in which the first promoter is inactive and the second promoter is active in a human cell.
- 18. (currently amended) The method of claim <u>14</u> 15, wherein a recombinant virus baculovirus is provided in which the first promoter is inactive and the second promoter is active in non-permissive insect cell.
- 19. (currently amended) The method of claim 18, wherein a recombinant <u>virus</u> <u>baculovirus</u> is provided in which the first promoter is inactive and the second promoter is active in a non-permissive *Drosophila* cell.
- 20. (currently amended) The method of claim 14 15, wherein a recombinant virus baculovirus is provided in which the first promoter is a viral polyhedrin promoter or a P10 promoter.

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21. (currently amended) The method of claim 14 15, wherein a recombinant virus baculovirus is provided in which the second promoter is a CMV promoter, a RSV promoter, a SV40 promoter, a heat shock protein promoter, an OPIE2 promoter, a MT promoter, or an actin 5C promoter.

22. (currently amended) A method for producing a protein <u>product</u> in a non-permissive cell, comprising:

providing a recombinant virus baculovirus that comprises eapable of infecting a nonpermissive cell, which virus includes: a first nucleic acid sequence encoding a detectable marker
operably linked to a first promoter, wherein the first promoter is active in a permissive host cell
culture and is inactive in a non-permissive cell; and a second nucleic acid sequence comprising
an exogenous nucleic acid sequence encoding a the protein product operably linked to a second
promoter, wherein the second promoter is active in the non-permissive cell and is inactive in the
permissive host cell culture;

infecting a the permissive host cell culture with the recombinant virus baculovirus; selecting a viral plaque by identifying expression of the detectable marker; amplifying the baculovirus virus by growing the baculovirus from the selected viral plaque; and

infecting a non-permissive cell with the amplified virus baculovirus, wherein the non-permissive cell thereby produces the protein product encoded by the exogenous nucleic acid sequence and wherein the non-permissive cell does not express the detectable marker.

- 23. (currently amended) The method of claim 22, further comprising the step of re-infecting the non-permissive cell with a recombinant virus baculovirus.
- 24. (canceled)
- 25. (currently amended) The method of claim 22 24, wherein the non-permissive cell infected is a mammalian cell.

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26. (currently amended) The method of claim 22 24, wherein the non-permissive cell infected is an insect cell.

- 27. (currently amended) The method of claim 22 24, wherein the non-permissive cell is infected in vitro.
- 28. (currently amended) The method of claim 22 24, wherein the non-permissive cell is infected in vivo.